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Award Number: DAMD17-01-C-0039

TITLE: Drug and Vaccine Evaluation in the Human-Aotus Plasmodium
Falciparum Model

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REPORT DATE: July 2002

TYPE OF REPORT: Annual

PREPARED FOR: U.S. Army Medical Research and Materiel Command
Fort Detrick, Maryland 21702-5012

DISTRIBUTION STATEMENT: Approved for Public Release;
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20020910 094

REPORT DOCUMENTATION PAGE			Form Approved OMB No. 074-0188	
<p>Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing this collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302, and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503</p>				
1. AGENCY USE ONLY (Leave blank)		2. REPORT DATE July 2002		3. REPORT TYPE AND DATES COVERED Annual (1 Jul 01 - 30 Jun 02)
4. TITLE AND SUBTITLE Drug and Vaccine Evaluation in the Human-Aotus Plasmodium Falciparum Model			5. FUNDING NUMBERS DAMD17-01-C-0039	
6. AUTHOR(S) Nicanor Obaldia III, D.V.M.				
7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES) Promed Trading, S.A. Panama, Panama 33126 Panama E-Mail: nobaldia@gorgas.gob.pa			8. PERFORMING ORGANIZATION REPORT NUMBER	
9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS(ES) U.S. Army Medical Research and Materiel Command Fort Detrick, Maryland 21702-5012			10. SPONSORING / MONITORING AGENCY REPORT NUMBER	
11. SUPPLEMENTARY NOTES				
12a. DISTRIBUTION / AVAILABILITY STATEMENT Approved for Public Release; Distribution Unlimited			12b. DISTRIBUTION CODE	
13. ABSTRACT (Maximum 200 Words) This report presents data: On the evaluation of antimalarial drugs. Experimental drugs GJ-237 (WR282650; BP20546) and GJ-QZ (WR282651; BP20537) were non toxic when given to Aotus orally at 20 mg/kg for three consecutive days and only transient changes on blood leukograms and liver enzyme ALT were detected. These drugs were not effective alone or in combination with Chloroquine when Aotus monkeys infected with <i>Plasmodium falciparum</i> FVO strain were treated for three consecutive days. Significant differences on Lymphocytes and Granulocytes counts were found between male and female Aotus when hematological and renal chemistry values of feral laboratory adapted Aotus monkeys were determined. Artelinic Acid/Lysine Salt (WR255663 BP; 23565) intravenously for three consecutive days at 2 and 8 mg/kg respectively suppressed or cleared <i>P. falciparum</i> FVO parasitemia in Aotus monkeys. Artesunic acid administered intravenously for three consecutive days at 35 mg/kg cleared and cured parasitemias of <i>P. falciparum</i> FVO in infected Aotus monkeys.				
14. SUBJECT TERMS Malaria, Plasmodium, Antimalarial Drugs, Vaccines, Aotus, Monkeys			15. NUMBER OF PAGES 38	
			16. PRICE CODE	
17. SECURITY CLASSIFICATION OF REPORT Unclassified	18. SECURITY CLASSIFICATION OF THIS PAGE Unclassified	19. SECURITY CLASSIFICATION OF ABSTRACT Unclassified	20. LIMITATION OF ABSTRACT Unlimited	

FOREWORD

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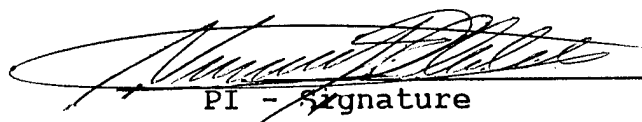
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INTRODUCTION:

Each year there are 300-500 million new infections and 2-5 million deaths attributable to malaria that occur primarily in countries in the tropics, particularly in sub-Saharan Africa (4). During the past 10-20 years the malaria problem has intensified in some parts of the world because parasites have developed resistance to drugs used for treatment and prevention; the anopheles mosquito, which transmits the parasite to humans, has developed resistance to insecticides, and control efforts have been reduced as resources have diminished in some developing countries (7).

The use of *Aotus lemurinus lemurinus* (Panamanian *Aotus* monkey), karyotypes VIII and IX (16) as a model to study malaria drug resistance and vaccine efficacy, have been ongoing at Gorgas Memorial Laboratory since 1976, due in part to the availability of this monkey in Panama (20), and also to the increasing drug resistance exhibited by the highly pathogenic *Plasmodium falciparum* parasites in Asia, Africa, and Latin America, and more recently *Plasmodium vivax* in the Melanesian and Indonesian archipelago (21). Previously, Schmidt (26,27) used the Colombian *Aotus* as the experimental host for antimalarial drug studies, but embargoes imposed by South American countries on the exportation of monkeys in the mid 1970's seriously restricted the use of *Aotus* for biomedical research in the United States, and in 1976 the project was transferred to Gorgas Memorial Laboratory where Panamanian *Aotus* were available for research. Five strains of *P. falciparum*, Vietnam Smith, Uganda Palo Alto, Vietnam Oak Knoll (FVO), Santa Lucia (5), and a C2A mefloquine resistant clone, and three strains of *P. vivax* Chesson (chloroquine sensitive), New Guinea AMRU-1 (chloroquine resistant) and Sal-1, have been adapted to Panamanian *Aotus*.

These strains exhibit diverse susceptibility and/or resistance to standard antimalarial agents. The course of untreated infections in Panamanian *Aotus* has been characterized and compared with that in *Aotus* of Colombia (25). Overall, the virulence of these strains was less in Panamanian than in Colombian owl monkeys, as indicated by lower mortality rates of Panamanian monkeys during the first 30 days of patency. Maximum parasitemias of the Vietnam Smith and Uganda Palo Alto strains were, however, significantly higher during the first 15 days of patency in Panamanian than in Colombian owl monkeys. These quantitative differences in infection parameters between Panamanian and Colombian owl monkeys have not invalidated the use of the former for evaluation of new antimalarial drugs.

Numerous candidate antimalarial drugs of diverse chemical classes have been evaluated against trophozoite-induced infections of one or more *P. falciparum* strains during the course of these contracts. In seeking alternatives to primaquine, two 8-aminoquinolines proved to be active against the blood stages of *P. falciparum* (2, 18). Desferrioxamine, an iron-specific chelating agent, was shown to suppress parasitemias of the virulent

Uganda Palo Alto strain of *P. falciparum* (23). The *in vitro* activity of two halogenated histidine analogs was not confirmed by evaluation against *P. falciparum* infections in owl monkeys (22).

Chloroquine-resistance of *P. falciparum* represents the greatest challenge in developing effective antimalarial drugs. Reversal of chloroquine-resistance in *P. falciparum*, *in vitro*, was achieved by the co-administration of verapamil (a calcium channel blocker) plus chloroquine (17). Other *in vitro* studies have shown that there is a significantly greater efflux of chloroquine from erythrocytes containing falciparum parasites resistant to chloroquine than from red cells parasitized by chloroquine-sensitive falciparum malaria (14). Calcium channel blockers appear to prevent this active efflux of chloroquine, thus allowing the drug to accumulate to parasitocidal levels.

Based upon the success of *in vitro* reversal of chloroquine-resistance, trials were initiated to determine if resistance could be reversed in *Aotus* infected with the chloroquine-resistant Vietnam Smith strain of *P. falciparum*. Six calcium channel blockers, or similarly acting drugs, were co-administered with chloroquine in diverse regimens. The desideratum of chloroquine-resistance reversal was administration of a single course of treatment, with parasite clearance and infection cure. Suppression of parasitemia was obtained during an initial course of treatment, but parasite clearance and cure occurred in some instances only after re-treatment. Such infection parameters were similar to those in monkeys with self-limited infections and cure could be attributed to acquired immunity.

Limited trials with desipramine, Norpramin, a tricyclic psychotropic drug, demonstrated the feasibility of reversing chloroquine-resistance *in vivo* (1). parasite clearance was obtained, but the infection was not cured.

Subsequently, *in vivo* reversal of chloroquine resistance was obtained with combinations of chloroquine plus chlorpromazine or prochlorperazine. Such reversal was exhibited by rapid suppression and clearance of parasitemia, resulting in infection cure without retreatment (15).

Evaluation of two oil-soluble derivatives of artemisinin, artemether and arteether, demonstrates that both possess similar activity to cure infections of a multi-drug resistant *P. falciparum* strain in *Aotus* (28).

Some strains of *P. vivax* from Melanesia and the Indonesian archipelago have demonstrated resistance to treatment with chloroquine (19, 24). Unlike chloroquine-resistant falciparum malaria, there exists no easy alternative to chloroquine-resistant strains of vivax malaria. Using WR 238605 alone or in combination with chloroquine in Panamanian *Aotus* monkeys it was demonstrated that WR238605 is an alternative treatment for chloroquine-resistant vivax malaria (21). The compound WR 238605 is a primaquine analog developed by the US Army as a better tolerated, more effective replacement for primaquine. Recent studies done at Gorgas Institute with Artemisinin derivative drugs developed by the U.S. Army such as Artelinic acid demonstrated its efficacy against the FVO strain of *P. falciparum* when administered orally to *Aotus l. lemurinus*.

Both the purpose and methods of approach of the present work remains essentially unchanged since 1976, viz to ascertain the antimalarial activity of drugs against *P. falciparum* and *P. vivax* in *Aotus*. The method of approach may vary on an ad hoc basis, such as administering a combination of drugs.

The long term goal of the second part of this project is to develop fully protective plasmid DNA vaccines that induce protective immune responses against the sporozoite, liver and erythrocytic stages of *P. falciparum*. If successful, it will establish, for the first time, that plasmid DNA vaccines can protect non-human primates, a critical step forward for the use of plasmid DNA vaccines in humans.

Vaccines are aimed at inducing immune responses that disrupt the complex cycle of the parasite at one more points: anti-sporozoite antibodies that prevent invasion of hepatocytes; cytotoxic T lymphocytes, cytokines, and antibodies that eliminate infected hepatocytes; antimerozoite antibodies that prevent invasion of erythrocytes; antibodies that neutralize parasite exoantigens known to induce harmful cytokine responses; antibodies that attack infected erythrocytes; cytokines that kill parasites within erythrocytes; and, anti-sexual stage antibodies that prevent the development of sporozoites in the mosquito.

Previous trials of malaria blood stage vaccines have shown that the Panamanian *Aotus*/*P. falciparum* model to be suitable for this purpose. (8-10).

Immunogenicity studies of a plasmid DNA vaccines encoding the circumsporozoite *P. yoelli* rodent malaria gene (PyCSP) in Panamanian *Aotus* monkeys demonstrated that the intradermal route of inoculation (ID) induces a higher level of antibodies than the intramuscular route (IM). Antibody levels induced in this manner reached a peak at week 9 and titers declined to 50% their peak value by week 14. When boosted at week 46 antibody levels increase 4 fold by week 49. This was comparable to antibodies generated with a Multiple Antigen synthetic peptide vaccine (MAP) delivered with an adjuvant (4)

We have used this immunization scheduled to test single or multi-gene DNA plasmid vaccines in *Aotus* monkeys. Additionally we have tested the ability of recombinant cytokines to enhance the immunogenicity and protective efficacy of the DNA vaccines. Preliminary using a small group of *Aotus l. lemurinus* (n=3) demonstrated partial, but incomplete, protection with a DNA vaccines for either AMA-1 or EBA-175 alone. These studies indicated that animals which received the vaccine candidates, had a short, but apparent significant delay in the onset of parasitemia {approximately 33% (1 of 3) self-cured, whereas none of the control animals did}. However, since the number of animals per group in each of these pilot studies were small, it was not possible to determined the absolute efficacy of these candidate vaccines, but these experiments suggested to the

investigators that further studies were warranted. MSP-1, when used as a protein/peptide vaccine formulation, provided protection from a *P. falciparum* infection in *Aotus* monkeys and we have demonstrated that, in mice and in Rhesus monkeys, the cytokine GM-CSF augmented both immunogenicity of a malaria DNA vaccine (personal communication. W. Weiss). We have now completed a pilot experiment to determine if *Aotus* Granulocyte-Macrophage-Colony Stimulating Factor (aGM-CSF) can augment immunogenicity and protective efficacy of a multi-gene erythrocytic vaccine.

In addition, synthetic oligodeoxynucleotides containing CpG motifs enhance immunogenicity of a peptide malaria vaccine when tested in Panamanian *Aotus* (11). Different vaccine formulations, routes and methods of administration with a comparable Hepatitis B Plasmid DNA vaccine were explored in Panamanian *Aotus* in order to elucidate the best route and methods of immunization for a plasmid DNA malaria vaccine (6). Further studies with single or multistage antigen plasmid DNA vaccines have been conducted or are in progress in Panamanian *Aotus* with variable results. Herein, we report partial protection obtained in *Aotus* monkeys immunized with either plasmid or recombinant protein in a primary and boosting immunization schedule using MSP1₄₂ as an antigen.

We have also tested the effect of prior *P. falciparum* infection on the immunogenicity of a DNA vaccine, obtaining partial protection in 67% of the monkeys (12). Also, evaluated in *Aotus* monkeys the characteristics of *P. falciparum*-induced anemia in two different experimental settings and hypothesis that a non-antibody/non-complement-mediated lysis of uninfected erythrocytes was the principal cause of anemia, and that bone marrow suppression and lysis of infected erythrocytes contributed to the anemia (13). In addition, we tested the hypothesis that a *P. falciparum* ligand, EBA-175 region II (RII), can be used as an immunogen in *Aotus* to induce antibodies that block the binding of RII to erythrocytes and thus inhibit parasite invasion of erythrocytes (29).

The purpose of this report is to: 1) Present data on the evaluation of potential antimalarial activity of drugs in the pre-clinical model of *Aotus l. lemurinus* (Panamanian night monkey) experimentally infected with *P. falciparum* or *P. vivax*, and 2) data on plasmid DNA and recombinant protein malaria vaccine experiments. These studies were supported by the U.S. Army and the U.S. Navy Malaria Programs.

BODY:

I. Experimental Methods

The first aim of this project is to evaluate the potential antimalarial activity of drugs, or combination thereof, in the preclinical model of *Aotus* experimentally infected with *P. falciparum* (or *P. vivax*). Specifically, the vertebrate host is *A. l. lemurinus*, the Panamanian night monkey. These animals are either feral, laboratory adapted or laboratory born. No naturally acquired, human plasmodium infection has been reported in *Aotus*. The Vietnam Smith/RE strain of *P. falciparum* was adapted to *Aotus* of Colombian origin in 1971 (26) and in Panamanian *Aotus* in 1976. (25). The course of untreated infections, essential for comparison with treated infections, has been documented in Panamanian *Aotus* (25). This plasmodium strain is resistant to maximally tolerated doses of chloroquine, pyrimethamine, and quinine (27).

To initiate an experiment, infected blood (with 2.5% sodium citrate as the anticoagulant) from an untreated *Aotus* was diluted appropriately in chilled saline (0.85%) or RPMI, such that each milliliter contained 5,000,000 parasites. This amount was inoculated into the saphenous vein of experimental and control monkeys.

Blood films, prepared and examined daily beginning on the first post-inoculation day, were stained with Giemsa. Parasitemias were evaluated as follows: negative, if no parasites were detected on a thick blood film after examination for at least 5 minutes; <10 parasites per cmm, if positive only on the thick blood film; parasite enumeration was by the Earle-Perez method and reported as the number of parasites per cmm. (3)

Blood films from untreated *Aotus*, serving as passage and/or control subjects, were prepared and examined daily during the primary patent period, and daily thereafter for at least three consecutive days after parasites could last be detected on thick blood films. When parasitemia had cleared, films were made and examined twice weekly until a total of 100 negative days had been recorded. If recrudescence occurred, blood films were obtained again on a daily basis.

Parasitemias were evaluated daily during the treatment period and until blood films were negative for at least seven consecutive days. The frequency of smearing was then reduced to two times per week (Monday and Thursdays or Tuesdays and Fridays). If no recrudescences occurred during a 100 day examination period, the infection was considered to have been cured.

Drug doses were calculated as mg base per kg of body weight. Stock solutions of water soluble compounds, at appropriate concentrations, were prepared with distilled water and stored at 8° C for the treatment period. If a compound was water insoluble, a suspension of the requisite amount of drug was prepared daily with 0.3% methylcellulose (in distilled water).

Oral administration of drugs was by gastric intubation with a 14 French catheter. The total volume of fluid administered, drug solution or suspension, and rinse was 14 ml.

Response to treatment was categorized as clearance and cure, clearance and recrudescence, or suppression without clearance. The day of clearance was defined as the first of three consecutive days in which the thick blood films were parasite negative. The day of recrudescence was the first of three consecutive days of positive thick blood films after a period of clearance. Suppression was defined as a transient decrease in the parasite count post-treatment without clearance.

The second objective of this project is to evaluate plasmid DNA vaccines against the blood and sporozoite stages of *P. falciparum* and against the blood stages of *P. vivax* in the Panamanian *Aotus* model. To this end we have evaluated single and multigene DNA vaccines of *both* *P. falciparum* and *P. vivax* with or without the addition of cytokines. The results of these experiments are detailed in results.

II Results

1. Toxicity of an oral route of administration of GJ-287 (WR282650; BP20546) and GJ-QZ (WR282651; BP20537) in *Aotus* monkeys.

In order to test the toxicity of an oral route of administration of GJ-287 (WR282650; BP20546) and GJ-QZ (WR282651; BP20537) in an *Aotus* monkey-model, in June 5, 2001, one (1) male and one (1) female *Aotus lemurinus lemurinus* malaria double cured monkeys weighing (845 and 900 gms), were administered 20 mg/kg of GJ-287 (WR282650; BP20546) or GJ-QZ (WR282651; BP20537) orally for three consecutive days respectively. The animals were monitored daily for signs such as: vomiting, depression, and anorexia and their weight was measured before treatment and twice a week after treatment. Blood was drawn from the femoral vein for CBC and Chemistry determinations such as Creatinine, Blood Urea Nitrogen and GPT before and during treatment and then twice a week for two weeks. No significant changes in body weight nor signs of toxicity were observed in the animals tested. In MN 12959 which received GJ-287 for three days a transient leukocytosis (WBC 49.87) with an absolute lymphocytosis (42.7%) was observed on the third day of treatment. Also during this period an increase in GPT (64.4 UI) was observed in this animal which then return to normal by day 10 post treatment (PT). In contrast, in MN 12960 which received GJ-QZ a leukopenia (WBC 9.71) with an absolute lymphocytosis (47.5%) was observed on the second day of treatment which then return to normal baseline values four days PT. A transient GPT increase on the first day of treatment (54.6 UI) was also present in this monkey.

2. To determine if the co-administration of GJ-287 (WR282650 BN; 20546) and GJ-QZ (WR282651 BP; 20537) alone or in combination with Chloroquine (WR1544 BM;AR20613) against infections of the FVO strain (CQR) of *Plasmodium falciparum* in Aotus monkeys reverse chloroquine resistance.

On June 11, 2001, each of sixteen (16) *Aotus l. lemurinus*, malaria naïve males and females weighing from (735-1046) grams, were divided into eight groups of two monkeys each and inoculated intravenously with 5×10^5 FVO strain of *P. falciparum* infected Erythrocytes. Blood films were obtained on the day after inoculation and continued daily for the duration of the experiment. When parasitemia reached 5,000 per cmm, oral treatment once a day for three days was administered as follows: Group 1. Received Chloroquine (WR1544 BM;AR20613) alone at 10 mg/kg once a day x three days. Group 2. Received GJ-287 (WR282650 BN; 20546) alone at 20 mg/kg once a day x three days. Group 3. Received GJ-QZ (WR282651 BP; 20537) alone at 20 mg/kg once a day x three days. Group 4. Received GJ-287 (WR282650 BN; 20546) at 20 mg/kg plus Chloroquine (WR1544 BM;AR20613) at 10 mg/kg once a day x three days. Group 5. Received GJ-QZ (WR282651 BP; 20537) at 20 mg/kg plus Chloroquine (WR1544 BM;AR20613) at 10 mg/kg once a day x three days. Group 6. Received GJ-287 (WR282650 BN; 20546) at 10 mg/kg plus Chloroquine (WR1544 BM;AR20613) at 10 mg/kg once a day x three days. Group 7. Received GJ-QZ (WR282651 BP; 20537) at 10 mg/kg plus Chloroquine (WR1544 BM;AR20613) at 10 mg/kg once a day x three days. Group 8. Served as Controls (no drug). As shown on table 42, 43 and 44 no effect over parasitemia was observed in the treated groups. One animal from group 4 and another one from group 5 died of malaria related complications on days 20 and 16 PI respectively.

3. Automated blood counts and renal function tests in feral laboratory adapted *Aotus lemurinus lemurinus* from Panamá.

Hematological values have been determined in the past in *Aotus lemurinus lemurinus* from Panamá (Karyotype VIII of IX) (Ma et al, 1978) using a manual system (Porter, 1969). Recently however, on May 14-17, 2001 these values were determined again at this laboratory as part of a Malaria protocol, using an automated hematological counter (MS4, France) and a chemistry analyzer (Reflotron, Merck). Herein, data on hematological and renal chemistry values are presented. Thirty (30) feral laboratory adapted (5-6 month in captivity) (fifteen (15) male and fifteen (15) female) adult *Aotus lemurinus lemurinus* monkeys and weighing between (700-950 grms) were bled from the femoral vein once to determine their blood and renal parameters which included CBC, creatininte and Blood Urea Nitrogen. Results are shown on table 45. No significant difference in CBC counts

between male and female monkeys were found except for lymphocytes ($p=0.05$) and granulocytes ($p=0.03$) differentials which were significantly different by one-tail student's T-test.

4. Efficacy of the intravenous administration of Artelinic acid/Lysine salt (WR255663 BP; 23565) against infections of *P. falciparum* FVO in Aotus monkeys.

Artelinic acid is currently under preclinical development for the treatment of multi-drug resistant and complicated malaria. The objective of this study was to determine the potential for use of IV artelinic acid using a new formulation in order to determine the optimal drug of choice for the treatment of severe malaria. On August 3, 2001, each of four (4) *Aotus l. lemurinus*, malaria naïve males and females weighing from (805-1038) grams were divided into 2 groups of two monkeys each and inoculated intravenously with 5×10^5 FVO strain of *P. falciparum* infected Erythrocytes. Blood films were obtained on the day after inoculation and continued daily for the duration of the experiment. Additionally, all animals were weighed, examined and a blood sample taken for CBC and chemistry analysis for serum creatinine and BUN determinations and serum samples stored at -70°C for further studies. When parasitemia reached 5,000 per cmm, intravenous treatment once a day for three days was administered as follows: Group 1 received at 2 mg/kg and Group 2 received 8 mg/kg of Artelinic Acid/Lysine salt (WR255663 BP; 23565) one control monkey received vehicle only. As shown in Table 5, 6 and 7 In Group 1, only one animal cleared its parasitemia on day 1 Post-treatment (PT), remaining negative for three days when recrudescence and was re-treated with the next highest dose at 8 mg/kg for three days, this time clearing on day 1 PT (17 PI) with two additional recrudescences on days 22 PI (22-28 PI) and 39 PI (39-42 PI) being treated with mefloquine at 20 mg/kg once orally on day 42 PI. The other animal suppressed its parasitemia for three days after treatment but had to be re-treated with the next highest dose on day 14 PI when its parasitemia reached 12,100 parasites x cmm, this time clearing on day 17 PI and remaining negative until day 22 PI when recrudescence (22-28 PI) and on day 39 PI (39-43 PI). Treatment with mefloquine at 20 mg/kg was instituted on day 43 PI. In group 2 both animals cleared on the second day PT but recrudescence on day 4 PT. One of these animals was retreated with 16 mg/kg of (WR255663 BP; 23565) on day 18 PI for three days, this time parasitemia cleared on day 21 PI and recrudescence on day 36 PI being treated with mefloquine on day 42 PI. The second animal of this group was re-treated with 16 mg/kg of (WR255663 BP; 23565) on day 21 PI for three days, clearing on day 24 PI and remaining negative until day 43 PI when it was treated with mefloquine. The control monkey was treated with 20 mg/kg of mefloquine on day 11PI when it reached 693,090 parasites x cmm. In conclusion, Aotus monkeys treated with Artelinic Acid/Lysine Salt

(WR255663 BP; 23565) intravenously for three consecutive days at 2 and 8 mg/kg respectively suppressed or cleared *P. falciparum* FVO parasitemia. Retreatment at 8 or 16 mg/kg cleared only. CBC determinations showed an increased in WBC counts with predominant granulocytosis in 3/4 treated animals. RBC, Hg concentration and platelets counts decreased in all animals during the experiment. In contrast, significant increases in ALT were observed in all. Weight lost of 7% body mass was found in 2/5 animals during the experiment.

5. Efficacy of the intravenous administration of Artelinic acid/Lysine salt (WR255663 BP; 23565) and Artesunic acid against infections of *P. falciparum* FVO in Aotus monkeys.

The objective of this study was to determine drug dosage levels of Artesunic Acid administered IV and compare it to Artelinic Acid/Lysine salt formulation. On 1 October 2001, each of ten (10) *Aotus l. lemurinus*, malaria naïve males and females weighing from (800-1096) grams were divided into 5 groups of two monkeys each and inoculated intravenously with 5×10^5 FVO strain of *P. falciparum* infected Erythrocytes. Blood films were obtained on the day after inoculation and continued daily for the duration of the experiment. Additionally, all animals were weighed, examined and a blood samples taken for CBC and chemistry analysis for serum ALT, creatinine and BUN determinations and serum samples stored at -70°C for further studies. When parasitemia reached 5,000 per cmm, treatment once a day for three days was administered intravenously as follows: Group 1 received Artelinic Acid/Lysine salt (WR255663 BP; 23565) at 32 mg/kg, Group 2 received Artesunic Acid IV at 2.2 mg/kg, Group 3 received Artesunic Acid IV at 8.8 mg/kg, Group 4 received Artesunic Acid at 35 mg/kg and group 5 received vehicle only. As shown on Tables 8, 9 and 10 In Group 1, one animal cleared its parasitemia on day 3 of treatment and the other one on the first day Post-Treatment (PT) remaining negative for 11 and 10 days respectively when recrudescence and were treated with mefloquine at 20 mg/kg once orally on day 24 PI. In Group 2 both animals cleared their parasitemias the first day PT but recrudescence on days 11 and 13 PT. They were retreated with Artesunic acid at 4.4 mg/kg for three days, this time one animal cleared on the third day of treatment and the other one on the second day PT. The first animal recrudescence on day 54 PI and self-cured on day 64 PI. The other animal recrudescence on day 60 PI and self-cured on day 63 PI. In group 3 both animals cleared on the first day PT and recrudescence on days 11 and 12 PT. Retreatment for three days with Artesunic acid at 17.6 mg/kg started on day 28 PI clearing on the second day PT and the other one on the third day of treatment. A second recrudescence occurred in one animal on day 68 PI and was treated with mefloquine at 20 mg/kg orally. The other animal cleared and cured. In group 4 both animals cleared their parasitemias on the third day of treatment. One animal cured its infection and the other one

recrudesce 12 days PT and was retreated on day 28 PI with Artesunic acid at 70 mg/kg, this time cured its infection on day 30 PI on the third day of treatment. Both control animals were treated with mefloquine at 20 mg/kg on day 12 PI when their parasitemias rised over 400,000 parasites x *ul*. CBC determinations showed an increased in WBC counts in most of the animal. RBC, Hg concentration and platelets counts decreased in all animals during the experiment. No increases were observed in creatinine or BUN but marked increases in ALT were observed in all animals. Weight lost up to 7% body mass was found in some of the animals during the experiment.

KEY RESEARCH ACCOMPLISHMENTS:

1. No significant changes in body weight nor signs of toxicity were observed in animals that received GJ-287 (WR282650; BP20546) or GJ-QZ (WR282651; BP20537) orally for three days. Only a transient leukocytosis with an absolute lymphocytosis or leukopenia and liver enzyme GPT increase were observed during the experiment.
2. No effect over parasitemia in Aotus infected with chloroquine resistant *P. falciparum* FVO, was observed when GJ-287 (WR282650 BN; 20546) or GJ-QZ (WR282651 BP; 20537) were administered orally alone or in combination with Chloroquine (WR1544 BM; AR20613).
3. Automated blood counts and renal function tests in feral laboratory adapted *Aotus lemurinus lemurinus* from Panamá demonstrated no significant difference in CBC counts between male and female monkeys except for lymphocytes ($p=0.05$) and granulocytes ($p=0.03$) differentials which were significantly different by one-tail student's T-test.
4. Artelinic Acid/Lysine Salt (WR255663 BP; 23565) intravenously for three consecutive days at 2 and 8 mg/kg respectively suppressed or cleared *P. falciparum* FVO parasitemia in Aotus monkeys.
5. Artesunic acid administered intravenously for three consecutive days at 35 mg/kg cleared and cured parasitemias of *P. falciparum* FVO in infected Aotus monkeys.

REPORTABLE OUTCOMES:

I. Manuscripts:

Jones TR, Gramzinski RA, Aguiar JC, Kim Lee Sim B, Narum DL, Furhmann SR, Kumar S, Obaldia NIII, Hoffman SL. 2002. Absence of antigenic competition in Aotus monkeys immunized with Plasmodium falciparum DNA vaccines delivered as a mixture. Vaccine 20(11-12):1675-80

Jones TR, Stroncek DF, Gozalo AS, Obaldia NIII, Andersen EM, Lucas C, Narum DL, Magill AJ, Sim BKL, Hoffman SL. 2002. Anemia in Parasite-and Recombinant Protein-Immunized Aotus Monkeys Infected with *P. falciparum*. Blood. Am J Trop Med Hyg. In Press

Sim KL, Narum DL, Liang H, Fuhrmann SR, Obaldia NIII, Gramzinski R, Aguiar J, Haynes DJ, Moch K, and Hoffman SL. 2001. Induction of biologically active antibodies in mice, rabbits, and monkeys by *Plasmodium falciparum* EBA-175 region II DNA vaccine. *Mol Med.* 7(4):247-254.

II. Presentations:

Jones TR, Gramzinski RA, Aguiar JC, Sim BKL, Narum DL, Fuhrmann Sr, Kumar S, Obaldia N, Hoffman SL. Absence of Antigenic Competition in Aotus Monkeys Immunized with *Plasmodium falciparum* DNA Vaccines Delivered as a Mixture. 50th Annual Meeting of The American Society for Tropical Medicine and Hygiene. Hilton, Atlanta, Georgia. November 11-15, 2001.

Ohrt C,....Obaldia N..... Status of Artelinic Acid development. 49th Annual Meeting of The American Society for Tropical Medicine and Hygiene. Westin Galleria & Oaks, Houston, Texas. October 29-November 2, 2000.

CONCLUSIONS:

1. Drugs GJ-287 (WR282650; BP20546) or GJ-QZ (WR282651; BP20537) were non toxic when given orally for three days to Aotus.
2. Drugs GJ-287 (WR282650 BN; 20546) or GJ-QZ (WR282651 BP; 20537) when administered orally alone or in combination with Chloroquine (WR1544 BM;AR20613) had no effect over parasitemia in *P. falciparum* FVO infected Aotus.
3. No significant differences in CBC counts or Renal function tests except for lymphocytes ($p = 0.05$) and granulocytes ($p = 0.03$) were found among male and female captive laboratory adapted Aotus monkeys.
4. Artelinic Acid/Lysine Salt (WR255663 BP; 23565) intravenously for three consecutive days at 2 and 8 mg/kg respectively suppressed or cleared *P. falciparum* FVO parasitemia in Aotus monkeys.
5. Artesunic acid administered intravenously for three consecutive days at 35 mg/kg cleared and cured parasitemias of *P. falciparum* FVO in infected Aotus monkeys

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TABLE 1

DETAILED ACTIVITY OF GJ-287* (WR282650 BN; 20546) AND GJ-QZ** (WR282651 BP;20537) WITH OR WITHOUT CHLOROQUINE*** (WR1544 BM;AR20613) AGAINST INFECTIONS OF *P. falciparum* FVO (CQR) IN AOTUS MONKEYS

RX INITIATED					PARASITEMIA PER cmmm x 10 ³								
MONKEY #	DAY P.I	DAY PAT.	MGKG	DAY PRE	DAY OF RX			DAY POST RX					
					RX	1	2	3	1	2	3	4	
													Days Neg.
13076	8	3	10***	0.13	22.5	3.3	180.7	652.3***	489	174.3	21.6	0	
13077	8	4	10***	0.2	17.5	20.4	309	899.9***	808	239.8	73.7	0	
13078	8	4	20*	0.01	2.7	2.2	46.5	220.4	365.4	901.4***	178	0	
13079	8	4	20*	0.15	8.5	2.9	108.7	418.2***	513	135.5	17.8	0	
13080	8	3	20**	0.11	6.9	1.1	196.6	176.6	733.2***	283.1	128.8	0	
13081	8	3	20**	0.01	14.4	2.8	211.4	795.7***	922.3	224.9	83.4	0	
13082	8	3	20*	0.01	1.34	0.18	13.8	5.9	45.3	217.4	203.1	(***)0	
13083	8	3	10***	0.01	26.7	7	108.7	197.8	290.6	1090.2***	707.7	0	
13084	8	2	20**	0.01	7.6	0.79	71.2	62.9	307.2	1203.4***	642.1	0	
13085	8	3	10***	0.01	3.9	0.68	20.7	1.34	44.2	124.5	196.3	(***)0	
13086	8	4	10*	0.18	16.8	1.4	295.9	371.4	1002.6***	625.5	163.9	0	
13087	8	3	10**	0.01	16.2	1.8	106.5	416***	238.5	31.2	25.3	0	
13088	8	3	10***	0.13	11	0.48	31	96.6	382	1697.2***	862.7	0	
13089	8	3	10**	0.01	18.5	20.2	168.7	223.4	549.6***	375	140	0	
13090	8	4	CONTROL	0.01	4.6	1.5	152.2	288.4	937.7***		135.5	0	
13091	8	3	CONTROL	0.01	3.1	0.42	46.5	85	61.4	671.9***	473.8	0	

***=Mefloquine 20 mg/kg

(***)=Mefloquine treated 20 mg/kg MN13082 day 15; died day 20 PI. MN13085 day 16; died day 20 PI.

TABLE 2

SUMMARY OF ACTIVITY OF GJ-287* (WR282650 BN; 20546) AND GJ-QZ** (WR282651 BP;20537) WITH OR WITHOUT CHLOROQUINE*** (WR1544 BM;AR20613) AGAINST INFECTIONS OF *P. falciparum* FVO (CQR) IN AOTUS MONKEYS

Monkey No.	Daily Dose x 3 mg/Kg	Response of parasitemia to Rx		Days from initial Rx to parasite Clearance	Days from final Rx to recrudescence	Notes No of days negative
		None	Suppressed	Cleared		
13076	10***	X				
13077	10***	X				
13078	20*	X				
13079	20*	X				
13080	20**	X				
13081	20**	X				
13082	20*	X				DIED day 20 PI
13083	10***	X				
13084	20**	X				
13085	20**	X				
13086	10*	X				
13087	10***	X				
13088	10**	X				
13089	10**	X				
13090	Control	X				
13091	Control	X				

TABLE 3

DETAILED PARASITEMIA OF GJ-287* (WR282850 BN; 20546) AND GJ-QZ** (WR282851 BP; 20537) WITH OR WITHOUT CHLOROQUINE*** (WR1544 BM; AR20613) AGAINST INFECTIONS OF *P. falciparum* FVO (CQR) IN AOTUS MONKEYS

P/DAY	GROUP	Parasitemia x cmm																			
		06/25/01	06/26/01	06/27/01	06/28/01	06/29/01	06/30/01	07/01/01	07/02/01	07/03/01	07/04/01	07/05/01	07/06/01	07/07/01	07/08/01	07/09/01	07/10/01	07/11/01	07/12/01	07/13/01	
13076	1	0	0	0.01	130	22500	3320	180750	652320***	489000	174330	21610	590	0.01	0	0	0	0	0	0	
13077	1	0	0.01	0.01	200	17510	20410	309000	899960***	808500	239890	73760	2310	0.01	0	0	0	0	0	0	
13078	2	0	0.01	0.01	0.01	2770	2220	46500	220460	365420	901470***	178060	34270	1140	0.01	0	0	0	0	0	
13079	2	0	0.01	0.01	150	8570	2930	108750	418270***	513000	135590	17880	17880	0.01	0	0	0	0	0	0	
13080	3	0	0	0.01	110	6900	1140	196680	176670	733290***	283100	128890	14900	0.01	0	0	0	0	0	0	
13081	3	0	0	0.01	0.01	14400	2840	211400	795770***	922310	224990	83440	3870	0.01	0	0	0	0	0	0	
13082	4	0	0	0.01	0.01	1340	180	13810	5980	45300	217440	203100	342020	459800***	235500	1050	0.01	DIED	0	0	
13083	4	0	0	0.01	0.01	26720	7020	108750	197810	290680	090220***	707750	41720	8940	0.01	0.01	0	0	0	0	
13084	5	0	0	0	0.01	7650	790	71250	62920	307290	203470***	642190	32780	4470	0.01	0.01	0	0	0	0	
13085	5	0	0	0.01	0.01	3980	680	20700	1340	44290	124500	196300	1015.5***	DIED	0.01	0.01	0	0	0	0	
13086	6	0	0.01	0.01	180	16800	1470	295960	371460	002640***	625500	163900	31290	790	0.01	0	0	0	0	0	
13087	6	0	0	0.01	0.01	16200	1870	106500	416010***	238580	31290	25330	0.01	0	0	0	0	0	0	0	
13088	7	0	0	0.01	130	11000	480	31000	96640	382030	697240***	862710	186.2	15650	730	0.01	0	0	0	0	
13089	7	0	0	0.01	0.01	18570	2020	168750	223480	549640***	375000	140060	15000	0.01	0.01	0	0	0	0	0	
13090	control	0	0.01	0.01	0.01	4600	1520	152250	288410	937710***	697320	135590	32010	0.01	0.01	0	0	0	0	0	
13091	control	0	0	0.01	0.01	3150	420	46500	85000	61410	671950***	473820	40230	0.01	0.01	0	0	0	0	0	

***= Treatment with Mefloquine 20 mg/kg

Treatment

TABLE 4 **AUTOMATED HEMATOLOGICAL AND RENAL CHEMISTRY VALUES OF
FERAL LABORATORY ADAPTED *Aotus l. lemurinus* MONKEYS FROM PANAMA**

MALES n=15

FEMALES n=15

	MEAN	STD	RANGE		MEAN	STD	RANGE	<i>p</i>
WBC X 10 ³	24.6	12.9	11.5-64.0		30.5	18.5	13.71-85.5	
LYM %	27.4	8.0	14.4-43.7		31.8	8.3	22.2-52.5	*0.05
MON %	20.1	4.7	11.8-28.7		22.5	6.6	8.9-33.0	
GRA %	52.6	11.2	35.2-69.2		45.6	12.0	25.6-64.4	*0.03
RBC X 10 ⁶	6.1	0.4	5.42-7.0		6.1	0.3	5.64-6.6	
MCV fl	87.0	3.1	81-91.8		86.9	3.2	80.2-90.6	
HCT %	52.9	3.3	47-58.9		52.7	2.9	47.7-56.9	
MCH pg	25.1	1.5	22.2-27.6		25.0	1.5	23-27.7	
RDW	29.1	1.3	27.431.4		28.8	1.4	26.7-30.7	
MCHC g/dl	8.2	0.4	7.4-8.9		8.4	0.3	7.9-8.8	
Hb g/dl	15.4	0.5	14.6-16.4		15.2	0.8	13.9-16.9	
PLT x 10 ³	528.9	122.8	338-771		513.3	118.0	343-764	
MPV fl	11.6	0.3	11.3-12.2		11.4	0.4	10.5-12.0	
Pct %	0.6	0.1	0.41-0.9		0.6	0.1	0.41-0.9	
PDW	10.1	0.8	8.0		10.2	0.8	8.9-11.6	
CREA mg/dl	0.5	0.0	0.5		0.5	0.0	0.5-0.6	
BUN mg/dl	10.0	0.0	10.0		10.9	2.3	10-17.8	

p= One Tail T-Test

TABLE 5

DETAILED ACTIVITY OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT (WR255663 BP; 23565)
AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

RX INITIATED				PARASITEMIA PER cmm x 10 ³								
ONKEY	DAY P.I	DAY PAT.	MGKG	DAY PRE	DAY OF RX			DAY POST RX				
RX					1	2	3	1	2	3	4 Days Neg.	
13019	8	6	2	0.9	36.3	13.5	0.33	0.01	0.01	0.01	12.1**	0
13024	8	6	2	0.51	29.7	12.3	0.01	0	0	0	0.01	3
13025	8	4	8	0.55	61.2	10	0.01	0.01	0	0	0.01	2
13028	8	6	8	0.53	49.5	5.5	0.26	0.01	0	0	0.01	2
13029	8	5	CONTROL	0.79	25.8	38.2	212.1	693*	311	79.2	26.5	0

*= Treated with mefloquine 40 mg/kg

**= Retreatment with next highest dose of Artelinic Acid

TABLE 6

SUMMARY OF ACTIVITY OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT (WR255663 BP; 23565)
AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

Monkey No.	Daily Dose x 3 mg/Kg	Responses of Parasitemia to Rx			Days from initial Rx to Parasite Clearance	Days from final Rx to parasite Clearance	Notes No. of days Negative
		None	Suppressed	Cleared			
13019	2	X					0
13024	2		X		4	1	3
13025	8		X		5	2	2
13028	8		X		5	2	2
13029	Control						0

TABLE 7

DETAILED PARASITEMIA OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT (WR255663 BP; 235665)
AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

		Parasites x cmm												
		DAY OF RX												
		08/04/01	08/05/01	08/06/01	08/07/01	08/08/01	08/09/01	08/10/01	08/11/01	08/12/01	08/13/01	08/14/01	08/15/01	
Monkey #	PI/DAY	2	3	4	5	6	7	8	9	10	11	12	13	
13019	1	0	0.01	0.01	0.01	550	900	36300	13500	330	0.01	0.01	0.01	
13024	1	0	0.01	0.01	0.01	510	510	29700	12370	0.01	0	0	0	
13025	2	0	0	0	0.01	680	550	61200	10000	0.01	0.01	0	0	
13028	2	0	0.01	0.01	0.01	420	530	49500	5500	260	0.01	0	0	
13029 control		0	0	0.01	0.01	240	790	25800	38250	212160	693090*	311060	79250	

*=Treated with mefloquine 40 mg/kg

**= Retreatment with next highest dose of Artelinic Acid

TABLE 7a

DETAILED PARASITEMIA OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT (WR255663 BP; 23565)
AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

		Parasites x cmm															
PI/DAY		08/16/01	08/17/01	08/18/01	08/19/01	08/20/01	08/21/01	08/22/01	08/23/01	08/24/01	08/25/01	08/26/01	08/27/01	08/28/01			
13019	12150**	640**	0.01**	0	0	0	0	0	0	0.01	0.01	0.01	0.01	190	180		
13024	0.01	1860	80940**	4500**	0.01**	0.01**	0	0	0	0.01	0.01	0.01	0.01	0.01	0		
13025	0.01	0.01	0.01	4840	1520	31200**	2400**	0.01**	0	0	0	0	0	0	0		
13028	0.01	0.01	0.01	0.01	0.01	260	1010	1500	10850**	2130**	0.01**	0	0	0	0		
13029	26500	670	0.01	0.01	0	0	0	0	0	0	0	0	0	0	0		

*=Treated with mefloquine 40 mg/kg

**= Retreatment with next highest dose of Artelinic Acid

TABLE 7b

DETAILED PARASITEMIA OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT (WR255663 BP; 235665)
AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

Parasites x cmm														
PI/DAY	08/29/01	08/30/01	08/31/01	09/01/01	09/02/01	09/03/01	09/04/01	09/05/01	09/06/01	09/07/01	09/08/01	09/09/01	09/10/01	
	27	28	29	30	31	32	33	34	35	36	37	38	39	
13019	310	0.01	0	0	0	0	0	0	0	0	0	0	0.01	
13024	0	0	0	0	0	0	0.01	0.01	460	620	1010	4050	7950	
13025	0	0	0	0	0	0	0	0	0	0.01	0.01	0.01	370	
13028	0	0	0	0	0	0	0	0	0	0	0	0	0	
13029	0	0	0	0	0	0	0	0	0	0	0	0	0	

*= Treated with mefloquine 40 mg/kg

**= Retreatment with next highest dose of Artelinic Acid

TABLE 7c

DETAILED PARASITEMIA OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT (WR255663 BP; 23565)
 AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

PI/DAY	Parasites x cmm													
	09/11/01	09/12/01	09/13/01	09/14/01	09/15/01	09/16/01	09/17/01	09/18/01	09/19/01	09/20/01	09/21/01	09/22/01	09/23/01	
13019	0.01	0.01	0.01	0.01*										
13024	21140	36750	23500*	1120										
13025	10630	8100	82500*	2210										
13028	0	0	0.01	0.01*										
13029	0	0	0	0										
*=Treated with mefloquine 40 mg/kg														
**= Retreatment with next highest dose of Artelinic Acid														

TABLE 8

DETAILED ACTIVITY OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT* (WR255663 BP; 23565)
AND ARTESUNIC ACID** AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

RX INITIATED				PARASITEMIA PER cmm x 10 ³									
MONKEY #	DAY P.I	DAY PAT.	MGKG	DAY PRE	DAY OF RX			DAY POST RX				Days Neg.	
					RX	1	2	3	1	2	3		4
13092	8	5	32*	3.4	5.8	1.1	0	0	0	0	0	0	11
13109	8	5	32*	1	4	1	0.01	0	0	0	0	0	10
13106	8	5	2.2**	2.1	7	0.33	0.01	0	0	0	0	0	11
13108	8	4	2.2**	1.1	2.8	0.22	0.01	0	0	0	0	0	13
13110	8	5	8.8**	1.6	5.7	0.44	0.01	0	0	0	0	0	10
13095	8	5	8.8**	1	3.5	0.79	0.01	0	0	0	0	0	11
13107	8	4	35**	1.5	3	0.01	0	0	0	0	0	0	11
13093	8	5	35**	1.4	7.5	0.46	0	0	0	0	0	0	100
13111	8	5	Vehicle AL	2.1	3.6	134.3	113.2	274	715.7***	267.7	76.5		
13094	8	5	Vehicle AS	1.9	4.1	87.3	47.3	312.3	247.6***	175.5	10.4		

***=Mefloquine 20 mg/kg

TABLE 9

SUMMARY OF ACTIVITY OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT (WR255663 BP; 23565)
AGAINST INFECTIONS OF P. FALCIPARUM FVO IN AOTUS MONKEYS

Monkey No.	Daily Dose x 3 mg/kg	Responses of Parasitemia to Rx			Days from initial		Days from final Rx to parasite Clearance	Notes No. of days Negative
					Rx to Parasite Clearance			
		None	Suppressed	Cleared				
13092	32*			X	3		-1	11
13109	32*			X	4		1	10
13106	2.2**			X	4		1	11
13108	2.2**			X	4		1	13
13110	8.8**			X	4		1	10
13095	8.8**			X	4		1	11
13107	35**			X	3		-1	11
13093	35**			X	3		-1	100
13111	Vehicle AL	X						
13094	Vehicle AS	X						

TABLE 10

DETAILED PARASITEMIA OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT* (WR255663 BP; 23565)
OR ARTESUNIC ACID IV** AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

		Parasitemia x cmm												
MONKEY #	GROUP PI/DAY	DAY OF Rx												
		10/03/01	10/04/01	10/05/01	10/06/01	10/07/01	10/08/01	10/09/01	10/10/01	10/11/01	10/12/01	10/13/01	10/14/01	
		2	3	4	5	6	7	8	9	10	11	12	13	
13092*	1	0	0	0.01	0.01	480	3480	5870	1120	0	0	0	0	
13109*	1	0	0	0.01	0.01	370	1080	4020	1030	0.01	0	0	0	
13106**	2	0	0	0.01	0.01	400	2160	7020	330	0.01	0	0	0	
13108**	2	0	0	0	0.01	0.01	1120	2860	220	0.01	0	0	0	
13110**	3	0	0	0.01	0.01	590	1610	5760	440	0.01	0	0	0	
13095**	3	0	0	0.01	0.01	170	1060	3590	790	0.01	0	0	0	
13107**	4	0	0	0	0.01	370	1520	3010	0.01	0	0	0	0	
13093**	4	0	0	0.01	0.01	310	1470	7500	460	0	0	0	0	
13111	Control AL	0	0	0.01	0.01	220	2180	3650	134390	113250	274070	715740***	267750	
13094	Control AS	0	0	0.01	0.01	590	1960	4180	87370	47310	312370	247640***	175550	

***=Mefloquine 20 mg/kg

****=Treated with next highest dose

TABLE 10a

DETAILED PARASITEMIA OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT* (WR255663 BP; 23565)
OR ARTESUNIC ACID IV** AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

Monkey #	GROUP P/DAY	Parasitemia x cmm														10/26/01
		10/15/01	10/16/01	10/17/01	10/18/01	10/19/01	10/20/01	10/21/01	10/22/01	10/23/01	10/24/01	10/25/01	10/26/01	10/27/01	10/28/01	
13092*	1	0	0	0	0	0	0	0	0.01	220	1030	1092***	1830			
13109*	1	0	0	0	0	0	0	0	0.01	500	2660	11180***	7200			
13106**	2	0	0	0	0	0	0	0	0	0.01	370	1140	11580			
13108**	2	0	0	0	0	0	0	0	0	0	0	0.01	460			
13110**	3	0	0	0	0	0	0	0	0.01	0.01	0.01	310	1390			
13095**	3	0	0	0	0	0	0	0	0	0.01	0.01	0.01	0.01			
13107**	4	0	0	0	0	0	0	0	0.01	0.01	0.01	420	680			
13093**	4	0	0	0	0	0	0	0	0	0	0	0	0			
13111 Control AL		76500	8160	0.01	0	0	0	0	0	0	0	0	0			
13094 Control AS		10470	1430	0.01	0.01	0	0	0	0	0	0	0	0			

***=Mefloquine 20 mg/kg

****=Treated with next highest dose

TABLE 10b

DETAILED PARASITEMIA OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT* (WR255663 BP; 23565)
OR ARTESUNIC ACID IV** AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

Monkey #	GROUP PI/DAY	Parasitemia x cmm													
		10/27/01 26	10/28/01 27	10/29/01 28	10/30/01 29	10/31/01 30	11/01/01 31	11/02/01 32	11/03/01 33	11/04/01 34	11/05/01 35	11/06/01 36			
13092*	1	0	0	0	0	0	0	0	0	0	0	0	0		
13109*	1	0	0	0	0	0	0	0	0	0	0	0	0		
13106**	2	12000	33220	40500****	4200****	0****	0	0	0	0	0	0	0		
13108**	2	13410	54360	119250****	36000****	3500****	0.01	0	0	0	0	0	0		
13110**	3	1043	40770	168000****	51750****	4000****	0.01	0	0	0	0	0	0		
13095**	3	200	840	1060****	0.01****	0****	0	0	0	0	0	0	0		
13107**	4	840	13500	26500****	920****	0****	0	0	0	0	0	0	0		
13093**	4	0	0	0	0	0	0	0	0	0	0	0	0		
13111 Control AL		0	0	0	0	0	0	0	0	0	0	0	0		
13094 Control AS		0	0	0	0	0	0	0	0	0	0	0	0		

***=Mefloquine 20 mg/kg

****= Treated with next highest dose

TABLE 10c

DETAILED PARASITEMIA OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT* (WR255663 BP; 23565)
OR ARTESUNIC ACID IV** AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

Monkey #	GROUP PI/DAY	Parasitemia x cmm											
		11/07/01 37	11/08/01 38	11/09/01 39	11/10/01 40	11/11/01 41	11/12/01 42	11/13/01 43	11/14/01 44	11/15/01 45	11/16/01 46	11/17/01 47	11/18/01 48
13092*	1	0	0	0	0	0	0	0	0	0	0	0	0
13109*	1	0	0	0	0	0	0	0	0	0	0	0	0
13106**	2	0	0	0.01	0	0	0	0	0	0	0	0	0
13108**	2	0	0	0	0	0	0	0	0	0	0	0	0
13110**	3	0	0	0	0	0	0	0	0	0	0	0	0
13095**	3	0	0	0	0	0	0	0	0	0	0	0	0
13107**	4	0	0	0	0	0	0	0	0	0	0	0	0
13093**	4	0	0	0.01	0	0	0	0	0	0	0	0	0
13111	Control AL	0	0	0	0	0	0	0	0	0	0	0	0
13094	Control AS	0	0	0	0	0	0	0	0	0	0	0	0

***= Mefloquine 20 mg/kg

****= Treated with next highest dose

TABLE 10d

DETAILED PARASITEMIA OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT* (WR255663 BP; 23565)
OR ARTESUNIC ACID IV** AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

Monkey #	GROUP	P/DAY	Parasitemia x cmm														11/30/01
			11/19/01	11/20/01	11/21/01	11/22/01	11/23/01	11/24/01	11/25/01	11/26/01	11/27/01	11/28/01	11/29/01	11/30/01	11/30/01	11/30/01	
			49	50	51	52	53	54	55	56	57	58	59	60	60	60	
13092*	1		0	0	0	0	0	0	0	0	0	0	0	0	0	0	
13109*	1		0	0	0	0	0	0	0	0	0	0	0	0	0	0	
13106**	2		0	0	0	0	0	0.01	0.01	460	700	510	440	590			
13108**	2		0	0	0	0	0	0	0	0	0	0	0	0.01			
13110**	3		0	0	0	0	0	0	0	0	0	0	0	0	0	0	
13095**	3		0	0	0	0	0	0	0	0	0	0	0	0	0	0	
13107**	4		0	0	0	0	0	0	0	0	0	0	0	0	0	0	
13093**	4		0	0	0	0	0	0	0	0	0	0	0	0	0	0	
13111	Control AL		0	0	0	0	0	0	0	0	0	0	0	0	0	0	
13094	Control AS		0	0	0	0	0	0	0	0	0	0	0	0	0	0	

***=Mefloquine 20 mg/kg

****=Treated with next highest dose

TABLE 10e

DETAILED PARASITEMIA OF INTRAVENOUS ADMINISTRATION OF ARTELINIC ACID/LYSINE SALT* (WR255663 BP; 23565)
OR ARTESUNIC ACID IV** AGAINST INFECTIONS OF *P. FALCIPARUM* FVO IN AOTUS MONKEYS

Parasitemia x cmm

Monkey #	GROUP PI/DAY	12/01/01 61	12/02/01 62	12/03/01 63	12/04/01 64	12/05/01 65	12/06/01 66
13092*	1	0	0	0	0	0	
13109*	1	0	0	0	0	0	
13106**	2	620	350	0.01	0	0	
13108**	2	0.01	0.01	0	0	0	
13110**	3	0	0	0	0	0	
13095**	3	0	0	0	0	0	
13107**	4	0	0	0	0	0	
13093**	4	0	0	0	0	0	
13111	Control AL	0	0	0	0	0	
13094	Control AS	0	0	0	0	0	

***=Mefloquine 20 mg/kg

****=Treated with next highest dose